Claims

1. An osteogenic device for implantation in a mammal, said device comprising:

a biocompatible, in vivo biodegradable matrix defining pores of a dimension sufficient to permit influx, proliferation and differentiation of migratory progenitor cells from the body of said mammal; and

a protein, produced by expression of recombinant DNA in a host cell, comprising one or more polypeptide chains, each of which has an amino acid sequence sufficiently duplicative of the sequence of COP-5 or COP-7 such that said protein is capable of inducing endochondral bone formation in association with said matrix when implanted in a mammal.

2. A device for implantation in a mammal, said device comprising:

a biocompatible, in vivo biodegradable matrix defining pores of a dimension sufficient to permit influx, proliferation and differentiation of migratory progenitor cells from the body of said mammal; and

a protein, produced by expression of recombinant DNA in a host cell, comprising one or more polypeptide chains, each of which has less than about 200 amino acids, in a sequence sufficiently duplicative of the sequence of

COP-5 or COP-7 such that said protein is capable of inducing cartilage formation in association with said matrix when implanted in a mammal.

3. The device of claim 1 or 2 wherein the sequence comprises:

4. The device of claim 1 or 2 wherein the sequence comprises:

5. The device of claim 1 or 2 wherein the sequence comprises:

10 20 30 40 50 CKRHPLYVDFRDVGWNDWIVAPPGYHAFYCHGECPFPLADHLNSTNHAIV RRRS K S S L QE VIS E FD Y E A AY MPESMKAS VI KE F E K I DN L N S Q ITK F P TL Q K A 60 70 80 90 100 QTLVNSVNPGKIPKACCVPTELSAISMLYLDENENVVLKNYQDMVVEGCGCR SI HAI SEQV EP A EQMNSLAI FFNDQDK I RK EE T DA H H RF T K DPV V Y N S S H RN RS N S K P E

wherein, in each position where more than one amino acid is shown, any one of the amino acids shown may be in that position.

6. The device of claim 1 or 2 wherein the sequence comprises:

10 20 30 40 50
LYVDFRDVGWNDWIVAPPGYHAFYCHGECPFPLADHLNSTNHAIV
K S S L QE VIS E FD Y E A AY MPESMKAS VI
F E K I DN L N S Q ITK F P TL
A S K

70 60 80 QTLVNSVNPGKIPKACCVPTELSAISMLYLDENENVVLKNYODMVVEGCGCR SI HAI SEQV EP A EQMNSLAI FFNDQDK I RK EE T DA H H RF K DPV V Y N S T S H RN RS S N K P E

wherein, in each position where more than one amino acid is shown, any one of the amino acids shown may be in that position.

7. The device of claim 1 or 2 wherein the sequence comprises:

1 10 20 30 40

Vg1 CKKRHLYVEFK-DVGWQNWVIAPQGYMANYCYGECPYPLTE
50 60 70

ILNGSN--H-AILQTLVHSIEPED-IPLPCCVPTKMSP
80 90 100

ISMLFYDNNDNVVLRHYENMAVDECGCR

8. The device of claim 1 or 2 wherein the sequence comprises:

DPP CRRHSLYVDFS-DVGWDDWIVAPLGYDAYYCHGKCPFPLAD

50 60 70

HFNSTN--H-AVVQTLVNNNNPGK-VPKACCVPTQLDS

80 90 100

VAMLYLNDQSTVVLKNYQEMTVVGCGCR

9. The device of claim 1 or 2 wherein the sequence comprises:

OP1 10 20 30 40

LYVSFR-DLGWQDWIIAPEGYAAYYCEGECAFPLNS
50 60 70

YMNATN--H-AIVQTLVHFINPET-VPKPCCAPTQLNA
80 90 100

ISVLYFDDSSNVILKKYRNMVVRACGCH

10.	The device of claim 1 or 2 wherein the
sequence	comprises:
OPl	-5 HQRQA 1 10 20 30 40 CKKHELYVSFR-DLGWQDWIIAPEGYAAYYCEGECAFPLNS 50 60 70 YMNATNH-AIVQTLVHFINPET-VPKPCCAPTQLNA 80 90 100 ISVLYFDDSSNVILKKYRNMVVRACGCH
11.	The device of claim 1 or 2 wherein the
sequence	comprises:
CBMP-2a	1 10 20 30 40 CKRHPLYVDFS-DVGWNDWIVAPPGYHAFYCHGECPFPLAD 50 60 70 HLNSTNH-AIVQTLVNSVNS-K-IPKACCVPTELSA 80 90 100 ISMLYLDENEKVVLKNYQDMVVEGCGCR
12.	The device of claim 1 or 2 wherein the
sequence	comprises:
CBMP-2b	1 10 20 30 40 CRRHSLYVDFS-DVGWNDWIVAPPGYQAFYCHGDCPFPLAD 50 60 70 HLNSTNH-AIVQTLVNSVNS-S-IPKACCVPTELSA 80 90 100 ISMLYLDEYDKVVLKNYQEMVVEGCGCR
13.	The device of claim 1 or 2 wherein the
sequence	comprises:
CBMP-3	1 10 20 30 40 CARRYLKVDFA-DIGWSEWIISPKSFDAYYCSGACQFPMPK 50 60 70 SLKPSNH-ATIQSIVRAVGVVPGIPEPCCVPEKMSS 80 90 100 LSILFFDENKNVVLKVYPNMTVESCACR

14. The device of claim 1 or 2 wherein the sequence comprises: 20 30 COPl LYVDFQRDVGWDDWIIAPVDFDAYYCSGACQFPSAD 60 HFNSTN--H-AVVQTLVNNMNPGK-VPKPCCVPTELSA 80 90 100 1SMLYLDENSTVVLKNYQEMTVVGCGCR The device of claim 1 or 2 wherein the 15. sequence comprises: 1 20 30 COP3 LYVDFQRDVGWDDWIVAPPGYQAFYCSGACQFPSAD 50 60 70 HFNSTN--H-AVVQTLVNNMNPGK-VPKPCCVPTELSA 80 90 100 **ISMLYLDENEKVVLKNYQEMVVEGCGCR** The device of claim 1 or 2 wherein the 16. sequence comprises: 20 30 1 COP4 LYVDFS-DVGWDDWIVAPPGYQAFYCSGACQFPSAD 50 60 70 HFNSTN--H-AVVQTLVNNMNPGK-VPKPCCVPTELSA 90 100 ISMLYLDENEKVVLKNYQEMVVEGCGCR The device of claim 1 or 2 wherein the 17. sequence comprises: 20 30 COP5 LYVDFS-DVGWDDWIVAPPGYQAFYCHGECPFPLAD 50 60 HFNSTN--H-AVVQTLVNSVNSKI--PKACCVPTELSA 80 90 ISMLYLDENEKVVLKNYQEMVVEGCGCR The device of claim 1 or 2 wherein the 18. sequence comprises: 10 20 30 40 COP7 LYVDFS-DVGWNDWIVAPPGYHAFYCHGECPFPLAD 50 60 70 HLNSTN--H-AVVQTLVNSVNSKI--PKACCVPTELSA 90 100 **ISMLYLDENEKVVLKNYQEMVVEGCGCR**

19. The device of claim 1 or 2 wherein the sequence comprises:

TO
PKHHSQRARKKNKN

1 10 20 30 40

COP16 CRRHSLYVDFS-DVGWNDWIVAPPGYQAFYCHGECPFPLAD
50 60 70
HFNSTN--H-AVVQTLVNSVNSKI--PKACCVPTELSA
80 90 100
ISMLYLDENEKVVLKNYQEMVVEGCGCR

- 20. The device of claim 1 or 2 wherein the osteogenics protein comprises a pair of separate polypeptide chains.
- 21. Osteogenic protein, produced by expression of recombinant DNA in a host cell, capable of inducing endochondral bone formation in association with a matrix when implanted in a mammal.
- 22. A protein, produced by expression of recombinant DNA in a host cell, comprising one or more polypeptide chains less than about 200 amino acids long in a sequence sufficiently duplicative of the sequence of COP-5 or COP-7 such that said protein is capable of inducing cartilage formation in association with a matrix when implanted in a mammal.
- 23. The osteogenic protein of claim 21 having an apparent molecular weight of about 30 kD when oxidized as determined by comparison to molecular weight standards in SDS-polyacrylamide gel.

- 24. The osteogenic protein of claim 23 further characterized by being glycosylated.
- 25. The osteogenic protein of claim 21 having an apparent molecular weight of about 27 kD as determined by comparison to molecular weight standards in SDS-polyacrylamide gel electrophoresis.
- 26. The protein of claim 22 or 25 further characterized by being unglycosylated.
- 27. The protein of claim 21 or 22 comprising a pair of separate polypeptide chains.
- 28. The protein of claim 21 or 22 comprising the amino acid sequences:

wherein each X independently represents an amino acid.

29. The protein of claim 21 or 22 comprising the amino acid sequences:

 30. The protein of claim 21 or 22 comprising the amino acid sequences:

10 20 30 50 CKRHPLYVDFRDVGWNDWIVAPPGYHAFYCHGECPFPLADHLNSTNHAIV RRRS K S S L QE VIS E FD Y E A AY MPESMKAS VI KE F E K I DN N S L Q ITK F P Α K 60 70 80 90 100 QTLVNSVNPGKIPKACCVPTELSAISMLYLDENENVVLKNYQDMVVEGCGCR SI HAI SEQV EP A EQMNSLAI FFNDQDK I RK EE T DA H H RF T S K DPV V Y N S H RN RS N S E P

wherein, in each position where more than one amino acid is shown, any one of the amino acids shown may be in that position.

31. The protein of claim 21 or 22 comprising the amino acid sequences:

10 20 30 40 50 LYVDFRDVGWNDWIVAPPGYHAFYCHGECPFPLADHLNSTNHAIV K S S L QE VIS E FD Y E A AY MPESMKAS VI N S FEKI Q ITK F P DN L TL A S K 70 60 80 90 QTLVNSVNPGKIPKACCVPTELSAISMLYLDENENVVLKNYQDMVVEGCGCR SI HAI SEQV EP A EQMNSLAI FFNDQDK I RK EE T DA H H RF T S K DPV V Y N S H RN RS N S K P Ε

wherein, in each position where more than one amino acid is shown, any one of the amino acids shown may be in that position.

32. The protein of claim 21 or 22 comprising the amino acid sequences:

1 10 20 30 40
Vgl CKKRHLYVEFK-DVGWQNWVIAPQGYMANYCYGECPYPLTE
50 60 70
ILNGSN--H-AILQTLVHSIEPED-IPLPCCVPTKMSP
80 90 100
ISMLFYDNNDNVVLRHYENMAVDECGCR

33. The protein of claim 21 or 22 comprising the amino acid sequences: 20 30 DPP CRRHSLYVDFS-DVGWDDWIVAPLGYDAYYCHGKCPFPLAD 50 60 HFNSTN--H-AVVQTLVNNNNPGK-VPKACCVPTQLDS 90 VAMLYLNDQSTVVLKNYQEMTVVGCGCR The protein of claim 21 or 22 comprising the amino acid sequence: 1 10 20 30 40 OP1 LYVSFR-DLGWQDWIIAPEGYAAYYCEGECAFPLNS 50 60 YMNATN--H-AIVQTLVHFINPET-VPKPCCAPTQLNA 90 **ISVLYFDDSSNVILKKYRNMVVRACGCH** The protein of claim 21 or 22 comprising the amino acid sequences: -5 HQRQA 20 30 40 OP1 CKKHELYVSFR-DLGWQDWIIAPEGYAAYYCEGECAFPLNS 50 60 YMNATN--H-AIVQTLVHFINPET-VPKPCCAPTQLNA 90 100 **ISVLYFDDSSNVILKKYRNMVVRACGCH** 36. The protein of claim 21 or 22 comprising the amino acid sequences: 20 30 CMP-2a CKRHPLYVDFS-DVGWNDWIVAPPGYHAFYCHGECPFPLAD 50 60 HLNSTN--H-AIVQTLVNSVNS-K-IPKACCVPTELSA 90 **ISMLYLDENEKVVLKNYQDMVVEGCGCR**

The protein of claim 21 or 22 comprising the amino acid sequences: 20 30 40 CBMP-2b CRRHSLYVDFS-DVGWNDWIVAPPGYQAFYCHGDCPFPLAD 50 60 HLNSTN--H-AIVQTLVNSVNS-S-IPKACCVPTELSA 90 100 **ISMLYLDEYDKVVLKNYQEMVVEGCGCR** 38. The protein of claim 21 or 22 comprising the amino acid sequences: 20 30 CBMP-3 CARRYLKVDFA-DIGWSEWIISPKSFDAYYCSGACOFPMPK 50 60 SLKPSN--H-ATIQSIVRAVGVVPGIPEPCCVPEKMSS 80 90 100 LSILFFDENKNVVLKVYPNMTVESCACR 39. The protein of claim 21 or 22 comprising the amino acid sequences: 10 20 30 COP1 LYVDFQRDVGWDDWIIAPVDFDAYYCSGACQFPSAD 50 60 70 HFNSTN--H-AVVQTLVNNMNPGK-VPKPCCVPTELSA 90 100 ISMLYLDENSTVVLKNYQEMTVVGCGCR 40. The protein of claim 21 or 22 comprising the amino acid sequences: 20 30 COP3 LYVDFQRDVGWDDWIVAPPGYQAFYCSGACQFPSAD 50 60 HFNSTN--H-AVVQTLVNNMNPGK-VPKPCCVPTELSA 90 **ISMLYLDENEKVVLKNYQEMVVEGCGCR** The protein of claim 21 or 22 comprising the amino acid sequences: 10 20 30 COP4 LYVDFS-DVGWDDWIVAPPGYQAFYCSGACQFPSAD

50

80 90 100
ISMLYLDENEKVVLKNYQEMVVEGCGCR

60

HFNSTN--H-AVVQTLVNNMNPGK-VPKPCCVPTELSA

42. The protein of claim 21 or 22 comprising the amino acid sequences:

1 10 20 30 40

COP5 LYVDFS-DVGWDDWIVAPPGYQAFYCHGECPFPLAD
50 60 70

HFNSTN--H-AVVQTLVNSVNSKI--PKACCVPTELSA
80 90 100

ISMLYLDENEKVVLKNYQEMVVEGCGCR

43. The protein of claim 21 or 22 comprising the amino acid sequences:

1 10 20 30 40

COP7 LYVDFS-DVGWNDWIVAPPGYHAFYCHGECPFPLAD
50 60 70

HLNSTN--H-AVVQTLVNSVNSKI--PKACCVPTELSA
80 90 100

ISMLYLDENEKVVLKNYQEMVVEGCGCR

44. The protein of claim 21 or 22 comprising the amino acid sequences:

-10
PKHHSSRARKKNKN

1 10 20 30 40

COP16 CRRHSLYVDFS-DVGWNDWIVAPPGYQAFYCHGECPFPLAD
50 60 70
HFNSTN--H-AVVQTLVNSVNSKI--PKACCVPTELSA
80 90 100
ISMLYLDENEKVVLKNYQEMVVEGCGCR

- 45. The protein of claim 21 or 22 comprising the product of expression of a DNA in a procaryotic cell.
- 46. A DNA sequence encoding an amino acid sequence sufficiently duplicative of that of the sequence encoded by the gene of Figure 1A_ such that said encoded sequence induces bone or cartilage formation when implanted in a mammal in association with a matrix.

- 47. The DNA of claim 46 encoding the same amino acid sequence as the gene set forth in Figure 1A.
- 48. The DNA sequence of claim 46 encoding:
- 1 10 20 30 40
 OP1 LYVSFR-DLGWQDWIIAPEGYAAYYCEGECAFPLNS
 50 60 70
 YMNATN--H-AIVQTLVHFINPET-VPKPCCAPTQLNA
 80 90 100
 ISVLYFDDSSNVILKKYRNMVVRACGCH
- 49. The DNA sequence of claim 46 encoding:

HQRQA

1 10 20 30 40

CKKHELYVSFR-DLGWQDWIIAPEGYAAYYCEGECAFPLNS

50 60 70

YMNATN--H-AIVQTLVHFINPET-VPKPCCAPTQLNA

80 90 100

ISVLYFDDSSNVILKKYRNMVVRACGCH

- 50. A cell line engineered to express the protein of claim 21 or 22.
- 51. The protein of claim 21 having a half maximum bone forming activity of about 20 25 ng per 25 mg of implant.
- 52. A biocompatible, in vivo biodegradable deglycosylated collagenous matrix defining pores of dimensions sufficient to permit influx, proliferation, and differentiation of migratory progenitor cells from the body of a mammal.
- 53. The matrix of claim 52 comprising close-packed particulate matter having a particle size within the range of 70-850 mm.

- 54. The matrix of claim 53 wherein said particulate matter has a particle size within the range of 70-420 mm.
- 55. The matrix of claim 52 defining a shape to span a non-union fracture in said mammal.
- 56. The matrix of claim 52 comprising demineralized, protein-extracted, deglycosylated, particulate xenogenic bone.
- 57. The matrix of claim 52 comprising a material selected from the group consisting of hydroxyapatite, tricalcium phosphate, polymers comprising lactic acid monomer units, polymers comprising glycolic acid monomer units, demineralized, guanidine-extracted, deglycosylated xenogenic bone, and mixtures thereof.
- 58. An osteogenic device for implantation in a mammal, said device comprising:

a biocompatible, in vivo biodegradable matrix defining pores of a dimension sufficient to permit influx, proliferation and differentiation of migratory progenitor cells from the body of said mammal; and

substantially pure osteogenic protein capable of inducing endochondral bone formation in said mammal disposed in said matrix and accessible to said cells.

- 59. The device of claim 1, 2, or 58 wherein said matrix comprises close-packed particulate matter having a particle size within the range of 70-850 mm.
- 60. The device of claim 1, 2, or 58 wherein said particulate matter has a particle size within the range of 70-420 mm.
- 61. The device of claim 1, 2, or 58 wherein said matrix comprises demineralized, protein-extracted, particulate, allogenic bone.
- 62. The device of claim 1, 2, or 58 wherein said matrix comprises a material selected from the group consisting of collagen, hydroxyapatite, tricalcium phosphate, polymers comprising lactic acid monomer units, polymers comprising glycolic acid monomer units, demineralized, guanidine-extracted allogenic bone, and mixtures thereof.
- 63. The device of claim 1, 2, or 58 wherein said matrix is shaped to span a non-union fracture in said mammal.
- 64. The device of claim 1, 2, or 58 disposed within the marrow cavity of allogenic bone.
- 65. The device of claim 1, 2, or 58 wherein said matrix comprises demineralized, protein extracted, particulate, deglycosylated xenogeneic bone.

- 66. The device of claim 65 wherein said matrix is treated with a protease.
- 67. The device of claim 58 wherein said osteogenic protein is unglycosylated.
- 68. The device of claim 67 wherein said osteogenic protein has an apparent molecular weight of about 27 kD when oxidized as determined by comparison to molecular weight standards in SDS-polyacrylamide gel electrophoresis.
- 69. The device of claim 58 wherein said osteogenic protein is glycosylated.
- 70. The device of claim 69 wherein said osteogenic protein has an apparent molecular weight of about 30 kD when oxidized as determined by comparison to molecular weight standards in SDS-polyacrylamide gel electrophoresis.
- 71. The device of claim 58 wherein said osteogenic protein comprises a pair of polypeptide chains.
- 72. The device of claim 71 wherein one chain of said pair of polypeptide chains has an apparent molecular weight of about 14 kD and the other has an apparent molecular weight of about 16 kD, both as determined after reduction by comparison to molecular weight standards in SDS-polyacrylamide gel electrophoresis.

73. The device of claim 71 wherein one chain of said pair of polypeptide chains has an apparent molecular weight of about 16 kD and the other has an apparent molecular weight of about 18 kD, both as determined after reduction by comparison to molecular weight standards in SDS-polyacrylamide gel electrophoresis.

74. The device of claim 58 wherein said osteogenic protein has the approximate amino acid composition set forth below:

Amino acid <u>residue</u>	Rel. no. res./molec.	Amino acid <u>residue</u>	Rel. no.
Aspartic acid	/ 22	Tyrosine	11
Asparagine		Valine	14
Glutamic acid	/ 24	Methionine	3
Glutamine		Cysteine	16
Serine	24	Isoleucine	15
Glycine	29	Leucine	15
Histidine 5		Proline	14
Arginine 13		Phenylalanine	e 7
Threonine 11		Tryptophan	ND
Alanine	18		
Lysine	12		

75. The device of claim 58 wherein said osteogenic protein comprises the amino acid sequence:

VPKPCCAPT

- 76. The device of claim 1 or 58 wherein the half maximum bone inducing activity of said protein is 0.8 to 1.0 ng per mg of said matrix.
- 77. A method of inducing local cartilage or bone formation in a mammal comprising the step of implanting the device of claim 1, 2, or 58 in said mammal at a locus accessible to migratory progenitor cells of said mammal.
- 78. A method of inducing endochondral bone formation in a mammal comprising the step of implanting the device of claim 1 or 58 in said mammal at a locus accessible to migratory progenitor cells of said mammal.
- 79. A method of inducing endochondral bone formation in a non-union fracture in a mammal comprising the step of implanting in the fracture in said mammal the device of claim 63.
- 80. Antibodies reactive with an epitope of the protein of claim 21 or 22.